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DB=DWPI; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L28	L27 and l26	62
<input type="checkbox"/>	L27	toxin	7426
<input type="checkbox"/>	L26	quinquestriatus or leiurus or scorpion	652

DB=JPAB; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L25	quinquestriatus or leiurus or scorpion	6
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DB=EPAB; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L24	L23 and l22	9
<input type="checkbox"/>	L23	toxin	1177
<input type="checkbox"/>	L22	quinquestriatus or leiurus or scorpion	29

DB=USOC; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L21	L20 same l19	0
<input type="checkbox"/>	L20	toxin	463
<input type="checkbox"/>	L19	quinquestriatus or leiurus or scorpion	28
<input type="checkbox"/>	L18	potassium or k+	105038

DB=PGPB; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L17	L15 and (potassium or k+)	109
<input type="checkbox"/>	L15	(L14 or l12) same l13	171
<input type="checkbox"/>	L14	scorpion	835
<input type="checkbox"/>	L13	toxin	22638
<input type="checkbox"/>	L12	quinquestriatus or leiurus	43

DB=USPT; PLUR=YES; OP=ADJ

<input type="checkbox"/>	L11	l7 and l6 not l9	61
<input type="checkbox"/>	L10	l7 and l3 not l8	33
<input type="checkbox"/>	L9	l7 same l6	34
<input type="checkbox"/>	L8	L7 same l3	3
<input type="checkbox"/>	L7	potassium or k+	282322
<input type="checkbox"/>	L6	L5 not l3	208
<input type="checkbox"/>	L5	l4 same l2	258
<input type="checkbox"/>	L4	scorpion	1172
<input type="checkbox"/>	L3	l1 same L2	53
<input type="checkbox"/>	L2	toxin	22671

L1 quinquestriatus or leiurus

79

END OF SEARCH HISTORY

S39	S38 AND S2	E30	1 AU=LEE JK-Y	Ref	Items Index-term
S40	338 D (sorted in duplicate order)	E31	77 AU=LEE JL	E1	15 AU=LEE JIAHN-SHING
S41	3201080 DNA OR GENE OR PLASMID	E32	4 AU=LEE JL C	E2	2 AU=LEE JIAHN-TE
S42	33 S39 AND S41	E33	2 AU=LEE JL JR	E3	0 AU=LEE JIAN
S43	33 ID (sorted in duplicate order)	E34	1 AU=LEE JL-C	E4	1 AU=LEE JIAN TAO
		E35	1005 AU=LEE JM	E5	1 AU=LEE JIAN-CHENG
		E36	4 AU=LEE JM F	E6	1 AU=LEE JIAN-FU
		E37	7 AU=LEE JM H	E7	13 AU=LEE JIAN-MING
		E38	2 AU=LEE JM J	E8	4 AU=LEE JIAN-WEI
		E39	1 AU=LEE JM JEANETTE	E9	1 AU=LEE JIAN GWU
		E40	3 AU=LEE JM JR	E10	4 AU=LEE JIAN-N-DER
		E41	1 AU=LEE JM L	E11	5 AU=LEE JIANN-FENG
		E42	1 AU=LEE JM TAMBLYN	E12	1 AU=LEE JIANN-GWU
		E43	2 AU=LEE JM-F	E13	1 AU=LEE JIANN-HSIUNG
		E44	2 AU=LEE JM MARSHALL	E14	3 AU=LEE JIANN-SHU
		E45	1 AU=LEE JM MARTIN		
		E46	1 AU=LEE JM MI		
		E47	26 AU=LEE JM MICHAEL		
		E48	162 AU=LEE J N		
S44	4131 AU='LEE J'	S45	1021 E35-E38,E41,E43		
				6/6/15 (item 15 from file: 5) 0005734218 BIOSIS NO.: 198784088367 ANALYSIS OF THE BLOCKING ACTIVITY OF CHARYBDOTOXIN HOMOLOGS AND IODINATED DERIVATIVES AGAINST CALCIUM-ACTIVATED POTASSIUM CHANNELS 1987	
				6/6/16 (item 16 from file: 5) 0007723137 BIOSIS NO.: 199191168028 CHARYBDOTOXIN-SENSITIVE CALCIUM ACTIVATED POTASSIUM CHANNEL IS NOT INVOLVED IN GLUCOSE-INDUCED ELECTRICAL ACTIVITY IN PANCREATIC BETA-CELLS 1991	
				6/6/17 (item 17 from file: 55) 09417323 PMID:1710672 Charybdotoxin -sensitive K(Ca) channel is not involved in glucose-induced electrical activity in pancreatic beta-cells. Jan 1991	
				6/6/18 (item 18 from file: 5) 0011348339 BIOSIS NO.: 198800143536 Consequence of the removal of evolutionary conserved disulfide bridges on particular cysteine spacing govern specific disulfide bond formation 1988	
				6/6/19 (item 19 from file: 5) 0007735556 BIOSIS NO.: 19919118447 DESIGN SYNTHESIS AND FUNCTIONAL EXPRESSION OF A GENE FOR CHARYBDOTOXIN A PEPTIDE BLOCKER OF POTASSIUM ION CHANNELS 1991	
				6/6/20 (item 20 from file: 55) 08571405 PMID: 2468961 Effect of some potassium channel blockers on contractile responses of the rabbit aorta. Feb 1989	
				6/6/21 (item 21 from file: 5) 0006633771 BIOSIS NO.: 198987058652 EFFECT OF SOME POTASSIUM CHANNEL BLOCKERS ON CONTRACTILE RESPONSES OF THE RABBIT AORTA 1989	
				6/6/22 (item 22 from file: 55) 08812992 PMID: 2531622 Effects of potassium channel toxins from Leishmania quinquestrigatus heteraeus venom on responses to cromakalim in rabbit blood vessels. Nov 1989	
				6/6/24 (item 24 from file: 55) 10238322 PMID: 7687466 Influence of protein surface charge on the bimolecular kinetics of a potassium channel peptide inhibitor. Jul 13 1993	
				6/6/25 (item 25 from file: 5) 0009404393 BIOSIS NO.: 199497425678 Evidence in support of a role for Ca(2+)-activated K+ channels in the hamster sperm acrosome reaction 1994	
				6/6/26 (item 26 from file: 55) 10570679 PMID: 7520055 Evidence in support of a role for Ca(2+)-activated K+ channels in the hamster sperm acrosome reaction 1994	
				6/6/27 (item 27 from file: 5) 0014215910 BIOSIS NO.: 200300174629 Functional analysis of an archaeabacterial voltage-dependent K+ channel. 2003	

- 6/6/28 (Item 28 from file: 5) 0006267342 BIOSIS NO.: 198886107263 PHARMACOLOGY OF POTASSIUM CHANNELS IN THE PLASMALEMMA OF THE GREEN ALGA CHARA-CORALLINA 1988
- 6/6/29 (Item 29 from file: 155) 07773735 PMID: 2433153 Identification of two toxins from scorpion (*Leurus quinquestrigatus*) venom which block distinct classes of calcium-activated potassium channel Dec 1 1986
- 6/6/30 (Item 30 from file: 155) 08722670 PMID: 2476127 Interactions between dendrotoxin, a blocker of voltage-dependent potassium channels, and charybdotoxin , a blocker of calcium-activated potassium channels, at binding sites on neuronal membranes. Aug 30 1989
- 6/6/31 (Item 31 from file: 5) 0006799478 BIOSIS NO.: 198988114593 INTERACTION BETWEEN DENDROTOXIN A BLOCKER OF VOLTAGE-DEPENDENT POTASSIUM CHANNELS AND CHARYBDOTOXIN A BLOCKER OF CALCIUM-ACTIVATED POTASSIUM CHANNELS AT BINDING SITES ON NEURONAL MEMBRANES 1989
- 6/6/32 (Item 32 from file: 5) 0011988561 BIOSIS NO.: 200400359350 Kbo1, a three disulfide bridges toxin from *Buthus occitanus turensanus* venom highly active on both SK and Kv channels 2004
- 6/6/33 (Item 33 from file: 5) 0006398088 BIOSIS NO.: 198926104699 LEURUS- QUINQUESTRIGATUS VENOM PEPTIDES THAT BLOCK BRAIN VOLTAGE-GATED AND CALCIUM-ACTIVATED POTASSIUM CHANNELS ALSO INHIBIT DENDROTOXIN BINDING TO SYNAPTIC MEMBRANES 1989
- 6/6/34 (Item 34 from file: 155) 10823421 PMID: 7819188 NMR sequential assignments and solution structure of charybdotoxin, a small scorpion toxin that blocks chloride channels. Jan 10 1995
- 6/6/35 (Item 35 from file: 5) 0006648827 BIOSIS NO.: 198598116660 NMR Sequential Assignments and Solution Structure of Charybdotoxin, a Small Scorpion Toxin That Blocks Chloride Channels 1995
- 6/6/36 (Item 36 from file: 155) 10805806 PMID: 7533051 Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Leiurus quinquestrigatus hebreus*. Nov 1994
- 6/6/37 (Item 37 from file: 5) 0008575215 BIOSIS NO.: 19859804048 Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Leiurus quinquestrigatus hebreus*. Nov 1994
- 6/6/38 (Item 38 from file: 155) 09390086 PMID: 1280139 Noriustoxin and leiumtoxin III, two homologous peptide toxins with binding properties to synaptosomal membrane. Sep 1992
- 6/6/39 (Item 39 from file: 5) 0008709107 BIOSIS NO.: 198395011373 Noriustoxin and leiumtoxin III, two homologous peptide toxins with binding properties to synaptosomal membrane potassium channels 1992
- 6/6/40 (Item 40 from file: 155) 06794986 PMID: 6197125 A study on the membrane depolarization of skeletal muscles caused by a scorpion toxin , sea anemone toxin II and crotamine and the interaction between toxins. Jul 1983
- 6/6/41 (Item 41 from file: 5) 0007309715 BIOSIS NO.: 198090094194 Polarized RUBIDIUM-86 EFFLUXES IN PRIMARY CULTURES OF RABBIT KIDNEY PROXIMAL CELLS ROLE OF CALCIUM AND HYPOTONICITY 1990
- 6/6/42 (Item 42 from file: 155) 09066202 PMID: 2165808 Polarized 86Rb+ effluxes in primary cultures of rabbit kidney proximal cells. role of calcium and hypotonicity. Jul 9 1990
- 6/6/43 (Item 43 from file: 5) 0005560519 BIOSIS NO.: 198783029410 PURIFICATION OF CHARYBDOTOXIN A SPECIFIC INHIBITOR OF THE HIGH-CONDUTTANCE CALCIUM-ACTIVATED POTASSIUM CHANNEL 1986
- 6/6/44 (Item 44 from file: 155) 07702384 PMID: 2429598 Purification of charybdotoxin , a specific inhibitor of the high-condutance Ca2+-activated K+ channel Nov 5 1986
- 6/6/45 (Item 45 from file: 155) 08257702 PMID: 2453055 Purification, sequencing, and model structure of charybdotoxin , a potent selective inhibitor of calcium-activated potassium channels. May 1988
- 6/6/46 (Item 46 from file: 5) 0006204047 BIOSIS NO.: 198886043968 PURIFICATION SEQUENCE AND MODEL STRUCTURE OF CHARYBDOTOXIN A POTENT SELECTIVE INHIBITOR OF CALCIUM-ACTIVATED POTASSIUM CHANNELS 1986
- 6/6/47 (Item 47 from file: 5) 0009198685 BIOSIS NO.: 19949718150 Solution structure of a core peptide derived from seyllatoxin 1994
- 6/6/48 (Item 48 from file: 155) 13206597 PMID: 1081954 Solution structure of potassium channel-inhibiting scorpion toxin Lq2. Mar 1 1999
- 6/6/49 (Item 49 from file: 5) 0008984542 BIOSIS NO.: 199497005827 Synthesis of charybdotoxin and of two N-terminal truncated analogues: Structural and functional characterization 1993
- 6/6/50 (Item 50 from file: 5) 0007639197 BIOSIS NO.: 199191022688 SYNTHESIS AND STRUCTURAL CHARACTERIZATION OF CHARYBDOTOXIN A POTENT PEPTIDYL INHIBITOR OF THE HIGH CONDUCTANCE CALCIUM-ACTIVATED POTASSIUM ION CHANNEL 1990
- 6/6/51 (Item 51 from file: 155) 09151531 PMID: 1639936 Synthesis and structural characterization of charybdotoxin , a potent peptidyl inhibitor of the high conductance Ca2(+)-activated K+ channel Nov 5 1990
- 6/6/52 (Item 52 from file: 5) 0010206293 BIOSIS NO.: 199698674126 Synthesis and structural characterization of arabyotoxin , a potassium channel blocker charybdotoxin 1996
- 6/6/53 (Item 53 from file: 5) 0009025088 BIOSIS NO.: 199497046373 Toxin pharmacology of the large-conductance Ca2+-activated K+ channel in the apical membrane of rabbit proximal convoluted tubule in primary culture. Oct 1993
- 6/6/54 (Item 54 from file: 155) 10398153 PMID: 7505914 Toxin pharmacology of the large-conductance Ca(2+)-activated K+ channel in the apical membrane of rabbit proximal convoluted tubule in primary culture. Oct 1993
- 6/6/55 (Item 55 from file: 155) 0974404035 PMID: 1373656 Toxin pharmacology of the ATP-induced hyperpolarization in Madin-Darby canine kidney cells. Mar 23 1992
- 6/6/56 (Item 56 from file: 5) 0008332816 BIOSIS NO.: 199294034657 TOXIN PHARMACOLOGY OF ATP-INDUCED HYPERPOLARIZATION IN MADIN-DARBY CANINE KIDNEY CELLS 1992
- 6/6/57 (Item 57 from file: 5) 0006361782 BIOSIS NO.: 198360901673 TOXINS IN THE CHARACTERIZATION OF POTASSIUM CHANNELS 1989
- 6/6/58 (Item 58 from file: 155) 10066261 PMID: 7678959 Toxin sensitivity of the calcium-dependent rubidium efflux in Madin-Darby canine kidney cells. Jan 29 1993
- 6/6/59 (Item 59 from file: 5) 0008801098 BIOSIS NO.: 19939510334 Toxin sensitivity of the calcium-dependent rubidium efflux in Madin-Darby canine kidney cells 1993
- 6/6/60 (Item 60 from file: 155) 08833309 PMID: 2600838 A voltage-dependent outward current with fast kinetics in single smooth muscle cells isolated from rabbit portal vein. May 1989
- 6/6/61 (Item 61 from file: 5) 0006708256 BIOSIS NO.: 198888023371 A VOLTAGE-DEPENDENT OUTWARD CURRENT WITH FAST KINETICS IN SINGLE SMOOTH MUSCLE CELLS ISOLATED FROM RABBIT PORTAL VEIN 1989
- 6/6/62 (Item 62 from file: 5) 0007742476 BIOSIS NO.: 1991125367 THREE-DIMENSIONAL STRUCTURE OF NATURAL CHARYBDOTOXIN IN AQUEOUS SOLUTION BY PROTON NMR CHARYBDOTOXIN POSSESSES A STRUCTURAL MOTIF FOUND IN OTHER SCORPION TOXINS 1991
- 6/7/2 (Item 2 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2006 Dialog. All its. reserv.
- 087580 PMID: 2477548 Analysis of the blocking activity of charybdotoxin homologs and iodinated derivatives against Ca2+-activated K+ channels.
- Lucchesi K, Ravindran A, Young H, Moczydlowski E Department of Pharmacology, Yale University School of Medicine, New Haven, Connecticut 06510. Journal of membrane biology (UNITED STATES) / Aug 1989, 109 (3) p269-81, ISSN 0022-2631 Journal Code: 0211301 Contract/Grant No.: AR38796; AR; NIAMS; HL38156; HL; NHBL Publishing Model Print Document type: Journal Article Languages: ENGLISH Main Citation Owner: NLM Record type: MEDLINE; Completed Microsequencing of the most abundant toxin, ChTX-Lq1, revealed identity with the 31-residue peptide previously sequenced by Gimenez-Gallego et al. [Gimenez-Gallego, G., et al., Proc. Natl. Acad. Sci. USA 85:3329-3333 (1988)]. Sequence data on the minor peptide, ChTX-Lq2, showed substantial homology to ChTX-Lq1 with differences observed at eight positions. These two charybdotoxin sequences, along with that of noxustoxin, define a distinct family of scorpion peptide toxins with activity against K+ channels. Both charybdotoxin homologs inhibited Ca2+-dependent K+ efflux from human erythrocytes with similar potency, K_{0.5} approximately 40 nM. In planar bilayer assays of single K(Ca) channels from rat muscle, ChTX-Lq1 and ChTX-Lq2 blocked with intrinsic K_ds of 1.3 and 43 nM, respectively, in the presence of 50 mM external KCl. A new application of dwell-time histogram analysis of single-channel blocking events was used to characterize the kinetic homogeneity of toxin samples and the blocking affinity of ChTX-Lq2

was the combined result of a faster dissociation rate and a slower association rate as compared to ChTX-Lq1. The blocking activity of two mono-iodinated derivatives of ChTX-Lq1 was also analyzed. Blocked dwell-time histograms of the iodinated peptides were characterized by predominately brief (0.2-2 sec) blocking events in comparison to the native toxin (20 sec). Histogram analysis revealed that mono-iodination of ChTX-Lq1 impairs blocking activity by adverse effects on both dissociation and association rate constants. Frequency density histograms of single channel blocking events provide a sensitive assay of toxin purity suitable for quantitating structure-activity relationships of charybotoxin derivatives.

Record Date Created: 1989/11/01 Record Date Completed: 1989/11/01

6776 (Item 6 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.

086775595 PMID: 2473920

Charybotoxin blocks both Ca-activated K channels and Ca-independent K channels in rat brain synaptosomes.

Schneider M J; Rogowski R S; Krueger B K; Blaustein M P

Department of Physiology, University of Maryland School of Medicine, Baltimore 21201.

FEBS Letters (NETHERLANDS) Jul 3 1989, 250 (2) p433-6, ISSN 0014-5793 Journal Code: 0155157

Contract/Grant No.: NS-16106; NS; NINDS; NS-16285; NS; NINDS; NS-20106; NS; NINDS; NS-20107

Document type: Journal Article Languages: ENGLISH Main Citation Owner: NLM Record type: MEDLINE; Completed

Charybotoxin (ChTX), a 4.3 kDa polypeptide toxin from the venom of the scorpion Leirus quinquestratus , blocks both a Ca-activated K channel (IC50 approximately 15 nM) and a Ca-independent voltage-gated K channel (IC50 approximately 40 nM) in rat brain synaptosomes. These results indicate that in this preparation ChTX is not specific for the Ca-activated K channel and suggest that there may be structural homology among the toxin-binding sites on various types of K channels.

Record Date Created: 1989/09/07 Record Date Completed: 1989/09/07

6779 (Item 9 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2006 Dialog. All rts. reserv.

11056434 PMID: 7513240

Charybotoxin and its effects on potassium channels.

Garcia M L; Kraus H G; Munujos P; Slaughter R S; Kaczorowski G J

Department of Membrane Biochemistry and Biophysics, Merck Research Laboratories, Rahway, New Jersey 07065, USA. American Journal of Physiology (UNITED STATES) Jul 1995, 268 (1 Pt 1) PC1-10, ISSN 0021-9361 Journal Code: 0370511 Publishing Model Print Document type: Journal Article; Review; Tutorial Languages: ENGLISH

Main Citation Owner: NLM Record type: MEDLINE; Completed

Over the last few years, a considerable amount of information has been obtained regarding K+ channels. Different areas of research have contributed to knowledge in this field. Charybotoxin (ChTX), a 37-amino acid peptide isolated from venom of the scorpion Leirus quinquestratus var. hebraeus, represents a remarkable tool for studying K+ channels. With its use, it has been possible to purify the high-conductance Ca(2+)-activated K+ (max-K) channel to homogeneity and determine the subunit composition of this channel. This has led to the discovery of an auxiliary beta-subunit that, when coexpressed with the pore-forming subunit, mSlo, alters the biophysical and pharmacological properties of this latter subunit. With the feasibility of producing large amounts of ChTX by recombinant techniques and the knowledge of the three-dimensional structure of the peptide, it has been possible to carry out site-directed mutagenesis studies and obtain a picture of the interaction surface of the toxin with two channels, maxi-K and Shaker, and to derive a picture of the complementary surface of the receptor in these two channels. Finally, ChTX, and the more selective K+ channel toxins that were subsequently discovered, have provided us with unique tools not only to determine the functional role that K+ channels play in target tissues but also to develop the molecular pharmacology of these channels. (76 Refs.)

Record Date Created: 1995/09/01 Record Date Completed: 1995/09/01

67711 (Item 11 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.

08859199 PMID: 2482078

Charybotoxin is a new member of the K+ channel toxin family that includes dendrotoxin 1 and mast cell degranulating peptide.

Schweitz H; Bidard J N; Maes P; Lazdunski M

Centre de Biochimie, Centre National de la Recherche Scientifique, Universite de Nice, France.

Biochemistry (UNITED STATES) Dec 12 1999, 28 (25) p9708-14, ISSN 0006-2960 Journal Code: 0370623

Publishing Model Print Document type: Journal Article Languages: ENGLISH Main Citation Owner: NLM Record type: MEDLINE; Completed

A polypeptide was identified in the venom of the scorpion Leirus quinquestratus hebraeus by its potency to inhibit the high-affinity binding of the radiolabeled snake venom toxin dendrotoxin 1 (125I-DTX1) to its receptor site. It has been purified, and its properties investigated by different techniques were found to be similar to those of MCD and DTX1, two polypeptide toxins active on a voltage-dependent K+ channel. However, its amino acid sequence was determined, and it was shown that this toxin is in fact charybotoxin (ChTX), a toxin classically used as a specific tool to block one class of Ca2+-activated K+ channels. ChTX, DTX1, and MCD are potent convulsants and are highly toxic when injected.

intracerebroventricularly in mice. Their toxicities correlate well with their affinities for their receptors in rat brain. These three structurally different toxins release [³H]GABA from preloaded synaptosomes, the efficiency order being DTX greater than ChTX greater than MCD. Both binding and cross-linking experiments of ChTX to rat brain membranes and to the purified MCD/DTX1 binding protein have shown that the alpha-subunit ($M_r = 78k\text{-}8k$) of the MCD/DTX1-sensitive K+ channel protein also contains the ChTX binding sites. Binding sites for DTX1, MCD, and ChTX are in negative allosteric interaction. Our results show that charybotoxin belongs to the family of toxins which already includes the dendrotoxins and MCD, which are blockers of voltage-sensitive K+ channels. ChTX is clearly not selective for Ca2+-activated K+ channel.

Record Date Created: 1990/03/05 Record Date Completed: 1990/03/05

67718 (Item 18 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

0011349339 BIOSIS NO.: 19980143566

Consequence of the removal of evolutionary conserved disulfide bridges on the structure and function of charybotoxin and evidence that particular cysteine spacing governs specific disulfide bond formation

AUTHOR: Drakopoulou Eugenia; Vizzanova Jean; Neyton Jacques; Aniort Vincent ; Bouet Fran^cois; Virelizier Henri; Menez Andre; Via Claudio (Reprint)

AUTHOR ADDRESS: CEA, Dep. Ingénierie Etudes Protéines, Serv. Phys. Exp. Analyse, Saclay, 91190 Gif-sur-Yvette, France**France

JOURNAL: Biochemistry 37 (5) p1292-1301 Feb. 3, 1998 1998 MEDIUM: print ISSN: 0006-2960 DOCUMENT TYPE: Article

RECORD TYPE: Abstract LANGUAGE: English

ABSTRACT: Scorpion toxins are miniglobular proteins containing a common structural motif formed by an alpha-helix on one face, an antiparallel beta-sheet on the opposite face, and three disulfide bonds making up most of its internal volume. We have investigated the role of these evolutionary conserved bonds by replacing each couple of bridged cysteine residues of the scorpion charybotoxin by a pair of nonbridging L-alpha-aminobutyric acid (Abu) residues. Three analogues were obtained by solid-phase synthesis, Chab I, Chab II, and Chab III, containing the Abu residues in positions 7 and 28, 13 and 33, 17 and 35, respectively. Circular dichroism analysis showed that the purified Chab II acquired a conformation similar to that of charybotoxin, while the Chab I and Chab III possess decreased native-like characteristics. All analogues block single high-conductance Ca2+-activated K+ channels from rat skeletal muscle inserted into planar lipid bilayers, but with different potencies. Chab II is the most active analogue (KD = 8.0 X 10-8 M), with a 9-fold lower affinity as compared to native charybotoxin . Chab I and Chab III have, respectively, 180- and 380-fold lower affinity. Therefore, the removal of evolutionary conserved disulfide bridges does not prevent the toxin to adopt a functional and presumably native-like structure. However, removal of one disulfide bond affects the yields of formation of correct pairing between the remaining cysteine residues, and only Chab I preserves the ability to form the native disulfide pairings with high efficiency. This is the only analogue to preserve partial spacings of three and one residue between the cysteines, which have been described to be thermodynamically disfavored disulfide bond formation between the cysteines (Zhang, R., and Snyder, G. H. (1989) J. Biol. Chem. 264, 18472-18479). Therefore, we conclude that the position of the cysteine residues in the sequence of charybotoxin , by disfavoring specific pairings and favoring others, may govern selective formation of specific disulfide bonds, thus, explaining the efficient folding properties of Chab I and of native charybotoxin . The structural properties of the Chab analogues and the discovered role of the cysteine spacings have interesting implications in protein design and engineering.

67719 (Item 19 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

0007735556 BIOSIS NO: 19911118447

DESIGN SYNTHESIS AND FUNCTIONAL EXPRESSION OF A GENE FOR CHARYBDOTOXIN A PEPTIDE BLOCKER OF POTASSIUM ION CHANNELS

AUTHOR: PARK C S; (Reprint); HAUSDORFF S F; MILLER C C

AUTHOR ADDRESS: HOWARD HUGHES MED INST GRADUATE DEP BIOCHEM, BRANDEIS UNIV, WALTHAM, MASS 02254, USA**USA

JOURNAL: Proceedings of the National Academy of Sciences of the United States of America 88 (6) p2046-2050 1991

ISSN: 0027-8424 DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: ENGLISH

ABSTRACT: A gene encoding charybotoxin (CTX), a K+ channel blocker from scorpion venom, was designed, synthesized, and expressed as a cleavable fusion protein in Escherichia coli. A sequence-specific protease, factor Xa, was used to cleave the fusion protein and thus release the toxin peptide. The recombinant toxin was purified, oxidized to form disulfide bonds, and treated to form N-terminal pyroglutamate. Recombinant CTX is identical to the native venom CTX with respect to high-performance liquid chromatography mobility, amino acid composition, and N-terminal modification. With single Ca2+-activated K+ channels as an assay system, recombinant CTX shows blocking and dissociation kinetics identical to the native venom toxin. The synthetic gene and high-level expression of functionally active CTX make it possible to study the fundamental mechanism of the toxin-ion channel interaction.

67722 (Item 22 from file: 155) DIALOG(R)File 155: MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.

08812992 PMID: 2531622

Effects of potassium channel toxins from Leurus quinquestratus hebraeus venom on responses to cromakalim in rabbit blood vessels. Strong P N; Wei S W; Beech D J; Hiestand P; Kocher H P

Jerry Lewis Muscle Research Centre, Department of Paediatrics and Neonatal Medicine, Royal Postgraduate Medical School, London.
British Journal of Pharmacology (ENGLAND) Nov 1989; 98 (3) p817-26, ISSN 0007-1188 .Journal Code: 7502536
Publishing Model Print Document type: Journal Article Languages: ENGLISH Main Citation Owner: NLM

Completed

1. The effects of fractionated *Lemurus quinquestratus* hebraeus venom on cromakalim-induced 86Rb⁺ efflux in rabbit aortic smooth muscle were examined. 2. Crude venom (0.1-10 micrograms ml⁻¹) produced a concentration-dependent decrease of 1 microm cromakalim-induced 86Rb⁺ response. The maximum blocking activity attainable was approximately 60%. 3. Fractionation of crude venom by gel permeation chromatography and subsequent chromatography on a cation ion-exchange column, produced two fractions (X and XI), active in the 86Rb⁺-blocking assay. 4. Fraction XII contained charybdotoxin (approximately 85% pure). After a final high performance liquid chromatography (h.p.l.c.) purification step, the purified toxin failed to inhibit the cromakalim-stimulated 86Rb⁺ efflux although it was a potent inhibitor of A23187-induced K⁺ flux in human erythrocytes, and the large conductance calcium-activated potassium channel in rabbit portal vein smooth muscle. 5. Subsequent purification of fraction X by h.p.l.c. yielded a minor peak which contained 86Rb⁺ blocking activity. This subfraction was also capable of inhibiting apamin-sensitive, angiotensin II-stimulated K⁺ flux in guinea-pig hepatocytes. 6. It is concluded that the potassium channel opened by cromakalim in rabbit aortic smooth muscle is not blocked by charybdotoxin but by another distinct toxin in the venom of *Lemurus quinquestratus* hebraeus.

Record Date Created: 198900125 Record Date Completed: 198900125

Identification of two toxins from scorpion (*Lemurus quinquestratus*) venom which block distinct classes of calcium-activated potassium channel
Castle N; Strong P N
FEBs letters (NETHERLANDS) Dec 1 1986; 209 (1) p17-21, ISSN 0014-5793 Journal Code: 0155157
Publishing Model Print Document type: Journal Article Languages: ENGLISH Main Citation Owner: NLM

Completed

Two polypeptide toxins from scorpion (*Lemurus quinquestratus*) venom which block distinct classes of calcium-activated potassium channels have been identified and partially purified. One toxin, at 50-100 ng/ml, blocks apamin-sensitive potassium fluxes in hepatocytes and inhibits [125I]monooctapeptide binding. The other, more basic, toxin blocks apamin-insensitive potassium fluxes in erythrocytes at 200 ng/ml and, to our knowledge, is the first toxin shown to block the erythrocyte calcium-activated potassium channel with high affinity. The possible co-identity of this latter toxin with charybdotoxin is discussed.

Record Date Created: 19870720 Record Date Completed: 19870720

67733 .(Item 33 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.
0006395608 BIOSIS NO.: 188936104699
LEMURUS QUINQUESTRATUS VENOM PEPTIDES THAT BLOCK BRAIN VOLTAGE-GATED AND CALCIUM-ACTIVATED POTASSIUM CHANNELS ALSO INHIBIT DENDROTOXIN BINDING TO SYNAPTIC MEMBRANES
AUTHOR: SOPENSEN R G (Reprint); SCHNEIDER M J; BLAUSTEIN M P
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JOURNAL: Biophysical Journal 55 (2 PART 2); p560A 1989 CONFERENCE/MEETING: THIRTY-THIRD ANNUAL MEETING OF THE BIOPHYSICAL SOCIETY, CINCINNATI, OHIO, USA, FEBRUARY 12-16, 1989. BIOPHYS J ISSN: 0006-3495 DOCUMENT TYPE: Meeting RECORD TYPE: Citation LANGUAGE: ENGLISH

67736 .(Item 36 from file: 5) DIALOG(R)File 555:MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.
10895806 PMID: 7533951
Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Lemurus quinquestratus* hebraeus.

Marshall D L; Vatrapour H; Harvey A L; Boyot P; Pinkasfeld S; Doljansky Y; Bouet F; Menez A
Department of Physiology and Pharmacology, University of Strathclyde, Glasgow, U.K.
Toxicon - official journal of the International Society on Toxicology (ENGLAND) Nov 1994; 32 (11) p1433-43, ISSN 0041-0101 .Journal Code: 1307333 Publishing Model Print .Document type: Journal Article Languages: ENGLISH Main Citation Owner: NLM Record type: MEDLINE; Completed
The scorpion venom *Lemurus quinquestratus* hebraeus was fractionated by chromatography in order to isolate toxins that affected binding of radiolabelled dendrotoxin to K⁺ channel proteins on synaptosomal membranes and that facilitated acetylcholine release in chick biventer cervicus nerve-muscle preparations. In addition to the previously characterized charybdotoxin , three toxins were isolated: 14-2, 15-1 and 18-2. Toxin 14-2 has a blocked N-terminus and because of low quantities, it has not been sequenced; 15-1 is a newly sequenced toxin of 36 residues with some overall homology to charybdotoxin and noxiustoxin; 18-2 is identical to charybdotoxin -2. The apparent Ki against dendrotoxin binding were: charybdotoxin , 3.8 nM, 14-2, 150 nM; 15-1, 50 nM, and 18-2, 0.25 nM. Toxin 14-2 (75 nM-1.5 microm) had a presynaptic

facilitation effect on neuromuscular preparations. Toxin 15-1 augmented responses to direct muscle stimulation, probably because it blocked Ca(2+)-activated K⁺ currents in muscle fibres. Toxin 18-2 (charybdotoxin -2) had a potent presynaptic facilitation action, with less effect on direct muscle stimulation. This contrasts with the relatively weak neuromuscular effects of the highly homologous charybdotoxin. On a Ca(2+)-activated K⁺ current in mouse motor nerve endings, charybdotoxin and toxin 18-2 produced maximal block at around 100 nM, whereas 15-1 was inactive at 300 nM. Charybdotoxin can increase quantal content, but this is more likely to result from block of voltage-dependent K⁺ channels than Ca(2+)-activated channels; the increase in transmitter release occurred in conditions in which little I_{KCa} would be present; higher concentration of charybdotoxin and longer exposure times were required to increase transmitter release than those needed to block I_{KCa}, and the facilitatory effects of charybdotoxin and toxin 18-2 correlated more with their effects on dendrotoxin binding than on block of I_{KCa}.

Record Date Created: 19980410 Record Date Completed: 19980410

67745 .(Item 45 from file: 155) DIALOG(R)File 155:MEDLINE(R) (c) format only 2005 Dialog. All rts. reserv.

08257702 PMID: 2453055
Purification, sequence, and model structure of charybdotoxin , a potent selective inhibitor of calcium-activated potassium channels.

Gimenez-Gallego G; Navia M A; Reuben J P; Katz G M; Kazazowski G J; Garcia M L

Department of Growth Factor Research, Merck Sharp & Dohme Research Laboratories, Rahway, NJ 07065.

Proceedings of the National Academy of Sciences of the United States of America (UNITED STATES) May 1988; 85 (10) p3229-33, ISSN 0027-8424 Journal Code: 7505876 Publishing Model Print Document type: Journal Article

Languages: ENGLISH Main Citation Owner: NLM Record type: MEDLINE; Completed

Charybdotoxin (ChTX), a protein present in the venom of the scorpion *Leurus quinquestratus* var. hebraeus, has been purified to homogeneity by a combination of ion-exchange and reversed-phase chromatography. Polycrylamide gel electrophoresis, amino acid analysis, and complete amino acid sequence determination of the pure protein reveal that it consists of a single polypeptide chain of 4.3 kDa. Purified ChTX is a potent and selective inhibitor of the approximately 220-pS Ca2+-activated K⁺ channel present in GH3 anterior pituitary cells and primary bovine aortic smooth muscle cells. The toxin reversibly blocks channel activity by interacting at the external pore of the channel protein with an apparent Kd of 2.1 nM. The primary structure of ChTX is similar to a number of neurotoxins of diverse origin, which suggests that ChTX is a member of a superfamily of proteins that modify ion-channel activities. On the basis of this similarity, the three-dimensional structure of ChTX has been modeled from the known crystal structure of alpha-bungarotoxin. These studies indicate that ChTX is useful as a probe of Ca2+-activated K⁺-channel function and suggest that the proposed tertiary structure of ChTX may provide insight into the mechanism of channel block. Record Date Completed: 19880622 Record Date Completed: 19880622

67752 .(Item 52 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

0010206283 BIOSIS NO.: 198698674126

Synthesis and structural characterisation of analogues of the potassium channel blocker charybdotoxin
AUTHOR: Dyke Timothy R; Duggan Brendan M; Pennington Michael W; Byrnes Michael E; Ken William R; Norton Raymond S (Reprint)
JOURNAL: Biophysical Research Inst, 381 Royal Parade, Parkville, VIC 3052, Australia**Australia
JOURNAL ADDRESS: NMR Lab., Biomolecular Research Inst, 381 Royal Parade, Parkville, VIC 3052, Australia**Australia
RECORD TYPE: Abstract LANGUAGE: English

ABSTRACT: Charybdotoxin is a 37-residue polypeptide toxin from scorpion venom, which acts by blocking voltage-gated and Ca2+-activated K⁺ channels. We have synthesized charybdotoxin and three mono-substituted analogues using an Fmoc-Ibu protocol. The Phe2 Iwdaw Tyr analogue was chosen to introduce a site for Tyr iodination which was distinct from the I⁺-channel binding surface, while the Glu-12 Iwdaw Glu and Arg-19 Iwdaw His analogues were studied to probe the roles of charged residues at these positions in the structure and activity of the toxin. The synthetic native molecule was equipotent with natural toxin in inhibiting the human erythrocyte Ca2+-dependent K⁺ channel. The affinities of all three analogues for the erythrocyte K⁺ channel were slightly reduced with the Arg-19 Iwdaw His analogue showing the greatest increase in IC-50 (2.3-fold). Two-dimensional 1H-NMR studies of these analogues showed that all three had structures very similar to those of the native molecule. The most significant perturbation was associated with the Glu-12 to Glu substitution, which appeared to destabilise the N-terminal half of the alpha-helix, possibly due to the weakening of an N-terminal helix capping interaction which is apparent from our NMR data. His-21 has a pKa more than one unit below the value for a non-interacting histidine. Possible reasons for this are that the imidazolium side chain is partly buried and is located near positively charged moieties. Thus, His-21 would be neutral at physiological pH, where charybdotoxin binds to the potassium channel.

67762 .(Item 62 from file: 5) DIALOG(R)File 5:Biosis Previews(R) (c) 2005 BIOSIS. All rts. reserv.

0007742476 BIOSIS NO.: 199191125367

THREE-DIMENSIONAL STRUCTURE OF NATURAL CHARYBDOTOXIN IN AQUEOUS SOLUTION BY PROTON NMR
CHARYBDOTOXIN POSSESSES A STRUCTURAL MOTIF FOUND IN OTHER SCORPION TOXINS
AUTHOR: BONTEMPS F (Reprint); ROUMESTAND C; BOYOT P; GILLQUIN B; DOLJANSKY Y; MENEZ A; TOMA F

- AUTHOR ADDRESS: SERV DE BIOCHIM DES PROTEINES, LAB D'INGENIERIE DES PROTEINES BAT 152, CEN-SACLAY, F-91191 GIF-SUR-YVETTE, FRANCE
- JOURNAL: European Journal of Biochemistry 1986 (1): p19-28 1991 ISSN: 0014-2956 DOCUMENT TYPE: Article
- RECORD TYPE: Abstract LANGUAGE: ENGLISH
- ABSTRACT: A 600-MHz proton NMR study of natural charybdotoxin , a toxin acting on K-channels, is reported. The analysis of NOEs and of backbone coupling constants showed the existence of an .alpha.-helix (residues 10-19) and of an antiparallel beta.-sheet in the 26 - 35 part. Three-dimensional structures were generated by distance geometry, using a set of 114 inter-residual calibrated constraints (63 sequential, 47 medium and long range, 4 hydrogen bonds) and 29 .PHI. angles. These structures show that charybdotoxin is composed of a beta.-sheet linked to an .alpha.-helix by two disulphide bridges and to an extended fragment by the third disulphide bridge. Comparison with the other known structures of long and short scorpion toxins shows that this structural motif is common to all these proteins.
- 9/6/1 (Item 1 from file: 5) 0007115178 BIOSIS NO.: 198089033069 ANALYSIS OF THE BLOCKING ACTIVITY OF CHARYBDOTOXIN HOMOLOGS AND IODINATED DERIVATIVES AGAINST CALCIUM-ACTIVATED POTASSIUM CHANNELS 1989
- 9/6/2 (Item 2 from file: 155) 08751830 PMID: 2477548 Analysis of the blocking activity of charybdotoxin homologs and iodinated derivatives against Ca²⁺-activated K+ channels. Aug 1989
- 9/6/3 (Item 3 from file: 155) 08675595 PMID: 2473920 Charybdotoxin blocks both Ca-activated K channels and Ca-independent voltage-gated K channels in rat brain synaptosomes. Jul 13 1989
- 9/6/4 (Item 4 from file: 155) 088559199 PMID: 2482078 Charybdotoxin is a new member of the K+ channel toxin family that includes dendrotoxin 1 and mast cell degranulating peptide. Dec 12 1989
- 9/6/5 (Item 5 from file: 5) 0007133033 BIOSIS NO.: 198089050924 CHARYBDOTOXIN IS A NEW MEMBER OF THE POTASSIUM CHANNEL TOXIN FAMILY THAT INCLUDES DENDROTOXIN 1 AND MAST CELL DEGRANULATING PEPTIDE 1989
- 9/6/6 (Item 6 from file: 5) 0006941512 BIOSIS NO.: 199038119403 CHARYBDOTOXIN IS A NEW MEMBER OF THE TOXIN FAMILY THAT INCLUDES DENDROTOXIN 1 AND MCD AND BLOCKS DENDROTOXIN-SENSITIVE VOLTAGE ACTIVATED POTASSIUM CHANNELS 1990
- 9/6/7 (Item 7 from file: 5) 0011349339 BIOSIS NO.: 199800143586 Consequence of the removal of evolutionarily conserved disulfide bridges on the structure and function of charybdotoxin and evidence that particular cysteine spacing govern specific disulfide bond formation 1998
- 9/6/8 (Item 8 from file: 155) 08571405 PMID: 24689361 Effect of some potassium channel blockers on contractile responses of the rabbit aorta. Feb 1989
- 9/6/9 (Item 9 from file: 5) 0006395808 BIOSIS NO.: 198936104699 LEIRUS: QUINQUESTRATUS VENOM PEPTIDES THAT BLOCK BRAIN VOLTAGE-GATED AND CALCIUM-ACTIVATED POTASSIUM CHANNELS ALSO INHIBIT DENDROTOXIN BINDING TO SYNAPTIC MEMBRANES 1989
- 9/6/10 (Item 10 from file: 155) 10823421 PMID: 7819188 NMR sequential assignments and solution structure of charybdotoxin, a small scorpion toxin that blocks chloride channels. Jan 10 1995
- 9/6/11 (Item 11 from file: 5) 0009564827 BIOSIS NO.: 199598116660 NMR Sequential Assignments and Solution Structures of Chirotoxin, a Small Scorpion Toxin That Blocks Chloride Channels 1995
- 9/6/12 (Item 12 from file: 155) 10895806 PMID: 7533951 Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Leirus quinquestriatus hebreus*. Nov 1994
- 9/6/13 (Item 13 from file: 5) 0009575215 BIOSIS NO.: 1995980403048 Neuromuscular effects of some potassium channel blocking toxins from the venom of the scorpion *Leirus quinquestriatus hebreus* 1994
- 9/6/14 (Item 14 from file: 155) 09939086 PMID: 1280139 Noxitoxin and leirutoxin III, two homologous peptide toxins with binding properties to synaptosomal membrane potassium channels 1992
- 9/6/15 (Item 15 from file: 5) 0008934542 BIOSIS NO.: 19949705527 Noxitoxin and leirutoxin III, two homologous peptide toxins with binding properties to synaptosomal membrane potassium channels 1992
- 9/6/16 (Item 16 from file: 5) 0008934542 BIOSIS NO.: 199698674126 Synthesis and structural characterisation of analogues of the potassium channel blocker charybdotoxin 1996
- 9/6/17 (Item 17 from file: 5) 0010206293 BIOSIS NO.: 199698674126 Synthesis and structural characterisation of analogues of the potassium channel blocker charybdotoxin 1996
- 9/6/18 (Item 18 from file: 5) 0009025088 BIOSIS NO.: 199497046537 Toxin pharmacology of the large-conductance Ca²⁺-activated K+ channel in the apical membrane of rabbit proximal convoluted tubule in primary culture 1993
- 12/6/1 (Item 1 from file: 155) 11322494 PMID: 8645186 A novel structural class of K+-channel blocking toxin from the scorpion *Pandinus imperator*. May 1 1996
- 13/6/1 (Item 1 from file: 155) 11322494 PMID: 8645186 A novel structural class of K+-channel blocking toxin from the scorpion *Pandinus imperator*. May 1 1996
- 13/6/2 (Item 1 from file: 5) 0010380507 BIOSIS NO.: 1996989014567 A novel structural class of K+-channel blocking toxin from the scorpion *Pandinus imperator* 1996
- 16/6/1 (Item 1 from file: 155) 12140757 PMID: 9438859 Solution structure and proposed binding mechanism of a novel potassium channel toxin kappa-conotoxin PVIIA. Dec 15 1997
- 16/6/2 (Item 1 from file: 5) 0011293829 BIOSIS NO.: 198800088076 Solution structure and proposed binding mechanism of a novel potassium channel toxin kappa-conotoxin PVIIA 1997
- 19/6/1 (Item 1 from file: 155) 10204659 PMID: 76856355 An activator of calcium-dependent potassium channels isolated from a medicinal herb. Jun 22 1993
- 19/6/2 (Item 2 from file: 155) 15271073 PMID: 15051409 Antigenic polymorphism of the "star" scorpion toxins able to block K+ channels. Mar 15 2004
- 19/6/3 (Item 3 from file: 5) 00095645783 BIOSIS NO.: 198783124674 BLOCKING AGENTS OF CALCIUM-ACTIVATED POTASSIUM CHANNELS IN CULTURED MEDULLARY THICK ASCENDING LIMB CELLS 1987
- 19/6/4 (Item 4 from file: 155) 07821071 PMID: 2435161 Blocking agents of Ca²⁺-activated K+ channels in cultured medullary thick ascending limb cells. Feb 1 1987
- 19/6/5 (Item 5 from file: 155) 17947714 PMID: 15695820 BmP9, a "big chain" scorpion peptide blocker of BK channels. Apr 15 2005
- 19/6/6 (Item 6 from file: 5) 0008848545 BIOSIS NO.: 199396012961 Calcium-activated potassium transport in erythrocytes: Comparison of binding and transport inhibition by scorpion toxins 1993
- 19/6/7 (Item 7 from file: 155) 10517036 PMID: 8297371 Chemical synthesis and structure-function studies of margatoxin, a potent inhibitor of voltage-dependent potassium channel in human T lymphocytes. Jan 28 1994
- 19/6/8 (Item 8 from file: 155) 08816237 PMID: 2483047 Characterization of high affinity binding sites for charybdotoxin in sacculenmal membranes from bovine aortic smooth muscle. Evidence for a direct association with the high conductance calcium-activated potassium channel. Dec 15 1989
- 19/6/9 (Item 9 from file: 155) 11411884 PMID: 8706835 Characterization of a new peptide from *Tityus serrulatus* scorpion venom which is a ligand of the apamin-binding site. Jul 15 1996
- 19/6/10 (Item 10 from file: 155) 16507366 PMID: 15188953 Computational simulations of interactions of scorpion toxins with the voltage-gated potassium ion channel Jun 2004
- 19/6/11 (Item 11 from file: 5) 0014960511 BIOSIS NO.: 200400331297 Computational simulations of interactions of scorpion toxins with the voltage-gated potassium ion channel 2004
- 19/6/12 (Item 12 from file: 155) 12172929 PMID: 9477955 Consequence of the removal of evolutionary conserved disulfide bridges on the structure and function of charybdotoxin and evidence that particular cysteine spacings govern specific disulfide bond formation. Feb 3 1998
- 19/6/13 (Item 13 from file: 155) 11091649 PMID: 7545007 Cross-linking of charybdotoxin to high-conductance calcium-activated potassium channels: identification of the covalently modified toxin residue. Aug 29 1995
- 19/6/14 (Item 14 from file: 155) 111965606 PMID: 8531201 Ca²⁺-activated K+ channels of human and rabbit erythrocytes display distinctive patterns of inhibition by venom peptide toxins. Sep 1995
- 19/6/15 (Item 15 from file: 155) 10145953 PMID: 7682555 Ca²⁺-activated K+ transport in erythrocytes. Comparison of binding and transport inhibition by scorpion toxins. Apr 25 1993
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Determination of the three-dimensional solution structure of noristoxin: analysis of structural differences with related short-chain scorpion toxins. Dec 26 1995

19618 (Item 18 from file: 155) 09906670 PMID: 1381959

Determination of the three-dimensional structure of ibenotoxin in solution by 1H nuclear magnetic resonance spectroscopy. Sep 8 1992

19619 (Item 19 from file: 155) 12195378 PMID: 9504384
Effect of *Thysanurus scutellatus* scorpion venom on the rabbit isolated corpus cavernosum and the involvement of NANC nitricergic nerve fibres. Feb 199819620 (Item 20 from file: 155) 13573819 PMID: 11155206
Effects of toxins P12 and P13 on human T lymphocyte Kv1.3 channels: the role of Glu7 and Lys24. Jan 1 200119621 (Item 21 from file: 155) 08469848 PMID: 2646066
An emerging pharmacology of peptide toxins targeted against potassium channels. Oct 198819622 (Item 22 from file: 155) 14309721 PMID: 12220678
Glycine 30 in ibenotoxin is a critical determinant of its specificity for maxi-K versus K(V) channels. Sep 11 200219623 (Item 23 from file: 155) 13705739 PMID: 11352729
Interaction of a toxin from the scorpion *Tityus serratus* with a cloned K+ channel from squid (sqKv1A). May 22 200119624 (Item 24 from file: 155) 17219256 PMID: 15165720
Kbot1, a three disulfide bridges toxin from *Buthus occitanus tunetanus* venom highly active on both SK and Kv channels. Apr 200419625 (Item 25 from file: 5) 000824656 BIOSIS NO.: 199293077547
KALIOTOXIN A NOVEL PEPTIDYL INHIBITOR OF NEURONAL BK-TYPE CALCIUM-ACTIVATED POTASSIUM CHANNELS CHARACTERIZED

FROM ANDROCTONUS-MAURETANICUS VENOM 1992

19626 (Item 26 from file: 5) 09629861 PMID: 1730708
Kaliotoxin, a novel peptidyl inhibitor of neuronal Bk-type Ca(2+)-activated K+ channels characterized from *Androctonus mauretanicus* mauretanicus venom. Jan 25 199219627 (Item 27 from file: 155) 10742600 PMID: 7524673
Kaliotoxin (1-37) shows structural differences with related potassium channel blockers. Nov 29 199419628 (Item 28 from file: 5) 00096 0999 BIOSIS NO.: 199593079832
Kaliotoxin (1-37) shows structural differences with related potassium channel blockers 199419629 (Item 29 from file: 155) 09853502 PMID: 1379069
Mechanism of ibenotoxin block of the large-conductance calcium-activated potassium channel from bovine aortic smooth muscle. Jul 28 199219630 (Item 30 from file: 155) 12969238 PMID: 10920011
Mechanisms of maurotoxin action on Shaker potassium channels. Aug 200019631 (Item 31 from file: 5) 00084 13156 BIOSIS NO.: 199294114987
MODE OF ACTION OF IBERIOTOXIN A POTENT BLOCKER OF THE LARGE CONDUCTANCE CALCIUM ACTIVATED POTASSIUM CHANNEL. 199219632 (Item 32 from file: 155) 09957833 PMID: 1384740
Mode of action of ibenotoxin, a potent blocker of the large conductance Ca(2+)-activated K+ channel. Aug 199219633 (Item 33 from file: 155) 14645123 PMID: 12660073
Molecular dynamics simulations of a K+ channel blocker: Tc1 toxin from *Thysanococcidae*. Jan 30 200319634 (Item 34 from file: 155) 141194806 PMID: 11864985
Mapping the binding site of a human ether-a-go-go-related gene-specific peptide toxin (EgGTX) to the channel's outer vestibule. May 10 200219635 (Item 35 from file: 5) 0013745123 BIOSIS NO.: 200200330634
Mapping the binding site of a human ether-a-go-go-related gene-specific peptide toxin (EgGTX) to the channel's outer vestibule 200219636 (Item 36 from file: 155) 09208395 PMID: 1702643
Mapping the receptor site for charybotoxin, a pore-blocking potassium channel inhibitor. Dec 199019637 (Item 37 from file: 155) 1343286 PMID: 10398897
A marine snail neurotoxin shares with scorpion toxins a convergent mechanism of blockade on the pore of voltage-gated K channels. Jul 199919638 (Item 38 from file: 5) 0012127174 BIOSIS NO.: 199900336834
A marine snail neurotoxin shares with scorpion toxins a convergent mechanism of blockade on the pore of voltage-gated K channels 199919639 (Item 39 from file: 155) 14633777 PMID: 12538890
Mutations and structural variability within toxins: implication for their use as scaffolds for protein engineering. Feb 200319640 (Item 40 from file: 5) 0014166498 BIOSIS NO.: 200300125608
Mutations and structural variability within toxins: Implication for their use as scaffolds for protein engineering. 200319641 (Item 41 from file: 155) 11833908 PMID: 9092804
NMR solution structure of a two-disulfide derivative of charybotoxin: 0 structural evidence for conservation of scorpion toxin alpha/beta motif19642 (Item 42 from file: 155) 11322494 BIOSIS NO.: 200000085730
A novel structural class of K+-channel blocking toxin from the scorpion *Pandinus imperator*. May 1 199619643 (Item 43 from file: 5) 0012367417 BIOSIS NO.: 200000085730
A point mutation in the maxi-K ctnb d5b forms a high affinity site for charybotoxin 199919644 (Item 44 from file: 155) 10274627 PMID: 8360176
Purification, characterization, and biosynthesis of mangatoxin, a component of *Centruroides margaritatus* venom that selectively inhibits voltage-dependent potassium channels. Sep 5 199319645 (Item 45 from file: 5) 0008972783 BIOSIS NO.: 199396137199
Purification, characterization, and biosynthesis of mangatoxin, a component of *Centruroides margaritatus* venom that selectively inhibits voltage-dependent potassium channels 199319646 (Item 46 from file: 155) 14164320 PMID: 11952787
Purification, characterization and biosynthesis of parabutoxin 3, a component of *Parabuthus transvaalensis* venom. Apr 200219647 (Item 47 from file: 5) 0013693181 BIOSIS NO.: 200200266892
Purification, characterization and biosynthesis of parabutoxin 3, a component of *Parabuthus transvaalensis* venom 200219648 (Item 48 from file: 155) 09024601 PMID: 1694715
Purification and characterization of a unique, potent, peptidyl probe for the high conductance calcium-activated potassium channel from venom of the scorpion *Buthus tamulus*. Jul 5 1990.19649 (Item 49 from file: 5) 0007278032 BIOSIS NO.: 199090063511
PURIFICATION AND CHARACTERIZATION OF A UNIQUE POTENT PEPTIDYL PROBE FOR THE HIGH CONDUCTANCE CALCIUM-ACTIVATED POTASSIUM CHANNEL FROM VENOM OF THE SCORPION *BUTHUS-TAMULUS* 199019650 (Item 50 from file: 155) 12065981 PMID: 9354615
Purification, characterization, and synthesis of three novel toxins from the Chinese scorpion *Buthus martensi*, which act on K+ channels. Nov 4 199719651 (Item 51 from file: 155) 12569895 PMID: 9891977
Purification and partial characterization of a 'short' insectotoxin-like peptide from the venom of the scorpion *Parabuthus schlecteri*. Dec 28 199819652 (Item 52 from file: 5) 0011824566 BIOSIS NO.: 199900084226
Purification and partial characterization of a 'short' insectotoxin-like peptide from the venom of the scorpion *Parabuthus schlecteri* 199819653 (Item 53 from file: 155) 10018942 PMID: 1467342
Progress in multidimensional NMR investigations of peptide and protein 3D structures in solution. From structure to functional aspects. Sep-Oct 199219654 (Item 54 from file: 5) 0008744162 BIOSIS NO.: 199395046428
Progress in multidimensional NMR investigations of peptide and protein 3-D structures in solution: From structure to functional aspects 199219655 (Item 55 from file: 155) 12054563 PMID: 9441593
[The rational evolution of scorpion toxins] Rational'naya evolyutsiya toksinov iz yadov skorpionov Sep 199719656 (Item 56 from file: 5) 0011277181 BIOSIS NO.: 199800071428
The rational evolution of scorpion toxins 199719657 (Item 57 from file: 155) 12019132 PMID: 9306273
Scorpion toxin block of the early K+-current (IK_A) in rat dorsal root ganglion neurones. Sep 1 199719658 (Item 58 from file: 5) 0011155973 BIOSIS NO.: 199799790333
Scorpion toxin block of the early K+-current (IK_A) in rat dorsal root ganglion neurones. Sep 1 199719659 (Item 59 from file: 155) 15269022 PMID: 15045653
Solution structure of BmBKTx1, a new BKCa1 channel blocker from the Chinese scorpion *Buthus martensi Karsch*. Apr 6 2004

- 196/60 (Item 60 from file: 5) 0014975010 BIOSIS NO.: 200400345799 Solution structure of BmBKTx1, a new BKCa channel blocker from the Chinese scorpion *Buthus martensi Karsch* 2004
- 196/61 (Item 61 from file: 155) 14028726 PMID: 11790849 Solution structure of a K⁺-channel blocker from the scorpion *Tityus cambridgei*. Feb 2002
- 196/62 (Item 62 from file: 155) 11808576 PMID: 9662103 Solution structure for Pandinus toxin K-alpha (PFTX-K alpha), a selective blocker of A-type potassium channels. Mar 11 1997
- 196/63 (Item 63 from file: 155) 12075374 PMID: 9655990 Solution structure of Tskapa, a charybdotoxin-like scorpion toxin from *Tityus serratus* with high affinity for apamin-sensitive Ca²⁺-activated K⁺ channels. Nov 1997
- 196/64 (Item 64 from file: 5) 0010328748 BIOSIS NO.: 1996081796381 Solution structure of SHK toxin, a novel potassium channel inhibitor from a sea anemone 1996
- 196/65 (Item 65 from file: 155) 11986450 PMID: 9252467 Sapecin B, a novel ly toxin, blocks macroscopic K⁺ currents in the GH3 rat pituitary cell line. Jul 1997
- 196/66 (Item 66 from file: 5) 0011067804 BIOSIS NO.: 199799701864 Sapecin B, a novellty toxin, blocks macroscopic K⁺ currents in the GH-3 rat pituitary cell line 1997
- 196/67 (Item 67 from file: 5) 0012839056 BIOSIS NO.: 200100010905 Structure determinants of scorpion toxin affinity: The charybdotoxin (alpha-KTx) family of K⁺-channel blocking peptides BOOK TITLE: Reviews of Physiology Biochemistry and Pharmacology 2000
- 196/68 (Item 68 from file: 155) 12908571 PMID: 10857399 Structural determinants of scorpion toxin affinity: the charybdotoxin (alpha-KTx) family of K⁺-channel blocking peptides. 2000
- 196/69 (Item 69 from file: 155) 12702762 PMID: 10707030 Structural and functional differences of two toxins from the scorpion *Pandinus imperator*. Mar 1 2000
- 196/70 (Item 70 from file: 5) 0012447874 BIOSIS NO.: 2000000166187 Structural and functional differences of two toxins from the scorpion *Pandinus imperator* 2000
- 196/71 (Item 71 from file: 5) 0010948956 BIOSIS NO.: 199799623016 Structural analysis of a two disulfide bridge analogue of a scorpion toxin 1997
- 196/72 (Item 72 from file: 155) 10377674 PMID: 8253752 Synthesis and characterization of kaiyotoxin. Is the 26-32 sequence essential for potassium channel recognition? Dec 15 1993
- 196/73 (Item 73 from file: 155) 10341798 PMID: 7693459 Synthesis of charybdotoxin and of two N-terminal truncated analogues. Structural and functional characterisation. Oct 1 1993
- 196/74 (Item 74 from file: 155) 11245554 PMID: 8547346 Synthesis and structural characterisation of analogues of the potassium channel blocker charybdotoxin . Jan 4 1996
- 196/75 (Item 75 from file: 155) 11180566 PMID: 7576559 Topology of the pore-region of a K⁺-channel revealed by the NMR-derived structures of scorpion toxins. Nov 1995
- 196/76 (Item 76 from file: 155) 10582194 PMID: 7514038 Tremorgenic indole alkaloids potently inhibit smooth muscle high-conductance calcium-activated potassium channels. May 17 1994
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- 24/6/2 (Item 2 from file: 5) 0010977935 BIOSIS NO.: 199799611995 Anti-insect toxin 5 (AaIT5) from *Androctonus australis* 1997
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- 24/6/4 (Item 4 from file: 155) 11165500 PMID: 8533143 Positive cooperativity among insecticidal scorpion neurotoxins. Aug 1995
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- 24/6/8 (Item 8 from file: 155) 10940776 PMID: 7722081 Direct effects of recombinant nuclear polyhedrosis viruses on selected nontarget organisms. Apr 1995
- 24/6/9 (Item 9 from file: 5) 0009776610 BIOSIS NO.: 19959824443 Direct effects of recombinant nuclear polyhedrosis viruses on selected nontarget organisms 1995
- 24/6/10 (Item 10 from file: 5) 00112496947 BIOSIS NO.: 20030025566 Effect of signal sequences and promoter on the speed of action of a genetically modified *Autographa californica* nuclear polyhedrovirus expressing the scorpion toxin Lqh112. 2003
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- 24/6/16 (Item 16 from file: 155) 14493200 PMID: 11782289 Isolation and characterization of a novel lepidopteran-selective toxin from the venom of South Indian red scorpion, *Mesobuthus tamulus*. 2001
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- 24/6/18 (Item 18 from file: 357) 0264969 DBR Accession No.: 2001-04723 New polynucleotides encoding scorpion venom potassium channel-agonist proteins for producing e.g. of insect-tolerant transgenic plants for controlling insect pest damage and parasitic worm infections - scorpion venom potassium channel-agonist protein genes useful for constructing transgenic plant with insect resistance 2000
- 24/6/19 (Item 19 from file: 357) 0264374 DBR Accession No.: 2001-04128 New isolated polynucleotide encoding a scorpion toxin for treating epilepsy, degenerative disorders such as Huntington's disease, and neuronal death following stroke, and for creating plants that are insect tolerant - transgenic plant construction with insect resistance and gene therapy 2000
- 24/6/20 (Item 20 from file: 5) 00086164579 BIOSIS NO.: 19934509567 Potential of baculoviruses expressing a scorpion toxin and an esterase in agriculture 1993
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Potential of baculoviruses expressing a scorpion toxin and an esterase in agriculture - use of recombinant baculovirus with a *Heliothis virescens* juvenile-hormone-esterase as an insect biological control agent (conference abstract) 1993

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Rapid purification and molecular modeling of AaIT peptides from venom of *Androctonus australis*. 1998

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Scorpion toxins 2004

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A SCORPION VENOM NEUROTOXIN PARALYTIC TO INSECTS THAT AFFECTS SODIUM CURRENT INACTIVATION PURIFICATION

PRIMARY STRUCTURE AND MODE OF ACTION 1990

24/626 (Item 26 from file: 155) 09075706 PMID: 2383565
A scorpion venom neurotoxin paralytic to insects that affects sodium current inactivation: purification, primary structure, and mode of action. Jun 26 1990

24/627 (Item 27 from file: 5) 0007645125 BIOSIS NO.: 199191028016
THE TOLERANCE OF LEPIDOPTEROUS LARVAE TO AN INSECT SELECTIVE NEUROTOXIN 1990

24/628 (Item 28 from file: 5) 0009069061 BIOSIS NO.: 199497090346
Variability among insect sodium channels revealed by selective neurotoxins 1994

24/629 (Item 29 from file: 5) 0010988996 BIOSIS NO.: 199799623056
Two novel short insectotoxins from the Asian scorpions *Buthus martensi* and *Buthus tamulus* 1997

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12339622 PMID: 9652392
A depressant insect-selective toxin analog from the venom of the scorpion Leirus quinquestratus hebraeus--purification and structure/function characterization.

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The scorpion-derived excitatory and depressant insect-selective polypeptide neurotoxins modify sodium conductance in insect membranes and differ greatly in their primary structures and symptoms induced in blow fly larvae. We report here the purification and characterization of a new insect selective toxin, LqhIT5. LqhIT5 is more similar to the excitatory toxins in its mode of action and the depressant toxins in its primary structure. This toxin is a single polypeptide composed of 61 amino acids that are cross linked by four disulfide bonds. When LqhIT5 is injected into blow fly larvae, a fast contraction paralysis occurs without depressant activity. No mammalian toxicity was detected by subcutaneous or intracranial injections of this toxin into mice. Sequence comparison of LqhIT5 and known depressant toxins shows a high degree of similarity among the amino acids located on the C-terminus of the toxins. However, there are some clear differences in the amino acids located close to the N-terminus of the toxins. By the aid of homology modeling, we demonstrated that these amino acids have the same orientation in the tertiary structure of the molecule and are exposed to the environment. The change in the mode of action of LqhIT5 (no depressant activity) by substitutions of a few amino acids located on a specific exposed area of the toxin shed a new light on the structure/function relationship of scorpion toxins. These results caution that similarity in the mechanism of action of scorpion toxins does not always follow from an overall similarity in sequence.
Record Date Created: 19980728 Record Date Completed: 19980728

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A depressant insect-selective toxin analog from the venom of the scorpion Leirus quinquestratus hebraeus--purification and structure/function characterization. May 15 1998

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A depressant insect-selective toxin analog from the venom of the scorpion Leirus quinquestratus hebraeus: Purification and structure/function characterization 1998

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Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin LqhIT2. 2003

26/66 (Item 6 from file: 357) 0313302 DBR Accession No.: 2003-14442
Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin LqhIT2 - scorpion venom expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

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A SCORPION VENOM NEUROTOXIN PARALYTIC TO INSECTS THAT AFFECTS SODIUM CURRENT INACTIVATION PURIFICATION PRIMARY STRUCTURE AND MODE OF ACTION 1990

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A scorpion venom neurotoxin paralytic to insects that affects sodium current inactivation: purification, primary structure, and mode of action. Jun 26 1990

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Variability among insect sodium channels revealed by selective neurotoxins 1994

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Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin LqhIT2 - scorpion venom expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

29/61 (Item 1 from file: 5) 0014419169 BIOSIS NO.: 20030377888
Scorpion toxins 2003

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Effect of signal sequences and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin LqhIT2 - scorpion venom expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

29/63 (Item 2 from file: 357) 0256837 DBR Accession No.: 2000-11327
New nucleic acid fragment encoding a scorpion toxin that is potassium channel-agonist, useful for creating transgenic plants that are more insect-tolerant - method is useful for producing transgenic plant with insect resistance 2000

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Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin LqhIT2. 2003

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Effect of signal sequences and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin LqhIT2 - scorpion venom expression in nuclear-polyhedrosis virus for biological control agent strain improvement 2003

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Scorpion toxins 2003

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The cDNA sequence of a depressant insect selective neurotoxin from the scorpion *Buthotus judaicus*. 1991

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THE cDNA SEQUENCE OF A DEPRESSANT INSECT SELECTIVE NEUROTOXIN FROM THE SCORPION BUTHOTUS-JUDAICUS 1991

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Cloning and characterization of a cDNA sequence encoding the precursor of a chirotoxin-like peptide from the Chinese scorpion *Buthus martensi* Karsch. Aug 2000

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Cloning and characterization of a cDNA sequence encoding the precursor of a chirotoxin-like peptide from the Chinese Scorpion *Buthus martensi* Karsch 2000

- 43665 (Item 5 from file: 5) 0014778747 BIOSIS NO.: 200400145408
Cytotoxic and apoptotic effects of scorpion *Leymus quinquestriatus* venom on 293T and C2C12 eukaryotic cell lines. 2003
- 43666 (Item 6 from file: 15) 10074904 PMID: 8431601
Depressant insect-selective neurotoxins from scorpion venom: chemistry, action, and gene cloning. 1993
- 43667 (Item 7 from file: 5) 0011914252 BIOSIS NO.: 199900174585
Dynamic diversification from a putative common ancestor of scorpion toxins affecting sodium, potassium, and chloride channels 1999
- 43668 (Item 8 from file: 5) 0007735566 BIOSIS NO.: 199191118447
DESIGN SYNTHESIS AND FUNCTIONAL EXPRESSION OF A GENE FOR CHARYBDOTOXINA PEPTIDE BLOCKER OF POTASSIUM CHANNELS 1991
- 43669 (Item 9 from file: 15) 12085528 PMID: 9395089
Influence of a NH2-terminal extension on the activity of KTx2, a K⁺ channel blocker purified from *Androctonus australis* scorpion venom. Nov 3 1997
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Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqht2. 2003
- 43671 (Item 11 from file: 357) 0313302 DBR Accession No.: 2003-14442
Effect of signal sequence and promoter on the speed of action of a genetically modified *Autographa californica* nucleopolyhedrovirus expressing the scorpion toxin Lqht2 - scorpion venom toxin expression in nuclear polyhedrosis virus for biological control agent strain improvement 2003
- 43672 (Item 12 from file: 155) 12401430 PMID: 9714546
Evidence for a new class of scorpion toxins active against K⁺-channels. Jul 24 1998
- 43673 (Item 13 from file: 155) 11211987 PMID: 7498537
Functional expression of an alpha anti-insect scorpion neurotoxin in insect cells and lepidopterous larvae. Dec 4 1995
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Functional expression of an alpha anti-insect scorpion neurotoxin in insect cells and lepidopterous larvae 1995
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Functional expression of an alpha anti-insect scorpion neurotoxin in insect cells and lepidopterous larvae - *Autographa californica* nuclear polyhedrosis virus vector allows insecticide expression in insect cell culture or larva 1995
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Genomic organization of three neurotoxins active no small conductance Ca²⁺-activated potassium channels from the scorpion *Buthus martensi Karsch* 1989
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Improved plant protective efficacy of a baculovirus using an early promoter to drive insect-specific neurotoxin expression 2005
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- 43681 (Item 21 from file: 5) 0012310167 BIOSIS NO.: 200000028480
Mortality and feeding of mid-stadium larvae of *Helicoverpa zea* and *Helicoverpa armigera* fed a wild strain or a recombinant strain of *Baculivirus* hebraeus expressing an insect-specific toxin of the scorpion *Leiurus quinquestriatus hebraeus* 1999
- 43682 (Item 22 from file: 155) 09709895 PMID: 1801321
Nucleotide sequence and structure analysis of a cDNA encoding an alpha insect toxin from the scorpion *Leiurus quinquestriatus hebraeus*. 1991
- 43683 (Item 23 from file: 5) 0008192856 BIOSIS NO.: 199293035717
NUCLEOTIDE SEQUENCE AND STRUCTURE ANALYSIS OF A cDNA ENCODING AN ALPHA INSECT TOXIN FROM THE SCORPION LEIURUS-QUINQUESTRIATUS -HEBRAEUS 1991
An 'Old World' scorpion beta-toxin that recognizes both insect and mammalian sodium channels. Jun 2003

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43666 (Item 26 from file: 5) 0013095637 BIOSIS NO.: 200100181376 Recombinant baculovirus insecticides 2000

43667 (Item 27 from file: 357) 01086334 DBR Accession No.: 90-11125 Recombinant DNA coding for insecticide toxin -including new scorpion venom toxin or spider venom toxin , and transgenic plant containing such DNA 1990

43668 (Item 28 from file: 357) 0148040 DBR Accession No.: 93-061092 Rapid isolation of full length cDNA clones by 'inverse PCR': purification of a scorpion cDNA family encoding alpha-neurotoxins - cDNA clone

43669 (Item 29 from file: 5) 0012648094 BIOSIS NO.: 200000368407 Scorpion neurotoxins: Structure/function relationships and application in agriculture 2000

43670 (Item 30 from file: 5) 0014419169 BIOSIS NO.: 200300377888 Scorpion toxins 2003

43671 (Item 31 from file: 5) 0012786805 BIOSIS NO.: 20000056518 Yields of occlusion bodies from *Heliothis virescens* (Lepidoptera: Noctuidae) and *Helicoverpa (Heliothis) zea* (Lepidoptera: Noctuidae) larvae fed wild or recombinant strains of baculoviruses 2000

43672 (Item 32 from file: 155) 14611132 PMID: 12467658 Three polymorphic genes encoding a depressant toxin from the Egyptian scorpion *Leiurus quinquestriatus* quinquestriatus .Jan 2003

43673 (Item 33 from file: 5) 0014146392 BIOSIS NO.: 200300105111 Three polymorphic genes encoding a depressant toxin from the Egyptian scorpion *Leiurus quinquestriatus* quinquestriatus .2003

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43675 (Item 1 from file: 357) 0264969 DBR Accession No.: 2001-04723 New polynucleotides encoding scorpion venom potassium channel-agonist proteins for producing e.g. of insect-tolerant transgenic plants for controlling insect pest damage and parasitic worm infections - scorpion venom potassium channel-agonist protein genes useful for constructing transgenic plant with insect resistance 2000

43676 (Item 2 from file: 357) 0264374 DBR Accession No.: 2001-04128 New isolated polynucleotide encoding a scorpion toxin for treating epilepsy, degenerative disorders such as Huntington's disease, and neuronal death following stroke, and for creating plants that are insect tolerant - transgenic plant construction with insect resistance and gene therapy 2000

43677 (Item 1 from file: 357) DIALOG(R)File 357-Derwent Biotech Res. (c) 2005 Thomson Derwent & ISI. All rts. reserv. 0264969 DBR Accession No.: 2001-04723 PATENT

New polynucleotides encoding scorpion venom potassium channel-agonist proteins for producing e.g. of insect-tolerant transgenic plants for controlling insect pest damage and parasitic worm infections - scorpion venom potassium channel-agonist protein genes useful for constructing transgenic plant with insect resistance

AUTHOR: Hermann R; Lee J M; Wong J F
CORPORATE SOURCE: DuPont 2000
PATENT NUMBER: WO 200078958 PATENT DATE: 2000-12-28 WPI ACCESSION NO.: 2001-071394 (2008)

PRIORITY APPLIC. NO.: US 140227 APPLIC. DATE: 19990622
NATIONAL APPLIC. NO.: WO 2000US17049 APPLIC. DATE: 20000621 LANGUAGE: English

ABSTRACT: An isolated polynucleotide (I) (e.g. DNA or RNA) is claimed. (I) contains a nucleotide sequence selected from a nucleotide sequence (III) of at least 81 nucleotides selected from 10 sequence of 171-213 (N1)-(N10), a nucleotide sequence (III) encoding a protein of at least 27 amino acids selected from 10 sequences of 56-70 amino acids (P1)-(P10), or a complement of (III) or (IV). Also claimed are: a chimeric gene or vector (III); a host cell (yeast, bacterium, plant) containing (I) or (II); a virus containing (I); a protein of at least 27 amino acids; a method of obtaining a nucleic acid fragment encoding a K-channel agonist; a recombinant baculovirus expression vector; and a method for testing the activity of a K-channel agonist against insects. (I) is useful for creating transgenic plants which express K-channel modifiers, useful as a means for controlling insect pest by producing insect tolerance. In the prevention or treatment of insect pest damage and parasitic worm infections in animals and humans, the invention may also find use in creating specific new pesticides and anthelmintic drugs that are also non-toxic to humans, pets and livestock. (50pp)